#### **AMENDMENTS TO THE CLAIMS**

What is claimed is:

1. (Currently Amended) A synthetic apolipoprotein-E mimicking polypeptide comprising an amino acid sequence

X-Y-Arg-Arg-Y-Y-X-X-Y-Y-Arg-Y-Y-Arg (SEQ ID NO: 210)

wherein X is glycine, threonine, serine or alanine,

wherein Y is a hydrophobic amino acid,

wherein the polypeptide comprises an acetyl group at the N-terminus and an amide group at the C-terminus, and

wherein the polypeptide consists of a single domain

wherein the polypeptide is capable of forming an amphipathic  $\alpha$  helical structure.

- 2. (Original) The polypeptide of claim 1, wherein Y is selected from the group consisting of phenylalanine, tyrosine, leucine, isoleucine, valine, and tryptophan.
- 3. (Currently Amended) The polypeptide of claim 1, wherein the polypeptide comprises from about 10 amino acids to about 30 14 amino acids to 18 amino acids in length.
- 4. (Currently Amended) The polypeptide of claim 1, wherein the polypeptide comprises a sequence of consecutive amino acids selected from the group of SEQ ID NOS: 2, 4, 5, 8, 10-11, 13, 18, 21, 110-121, 127, 129, 131, 133, 137, 141, 145, 150, 155-160, 167, 168, 194-196, and 203-204 of SEQ ID NO:5.
- 5. (Original) The polypeptide of claim 1, wherein the polypeptide comprises the sequence Gly-Ile-Arg-Arg-Phe-Leu-Gly-Ser-Ile-Trp-Arg-Phe-Ile-Arg-Ala-Phe-Tyr-Gly

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### (SEQ ID NO:5).

- 6. (Original) The polypeptide of claim 1, which is a recombinant polypeptide.
- 7. (Canceled)
- 8. (Canceled)
- 9. (Withdrawn) An isolated nucleic acid encoding the polypeptide of claim 1.
- 10. (Withdrawn) The nucleic acid of claim 9, wherein the nucleic acid comprises DNA, RNA and/or cDNA.
- 11. (Withdrawn) A vector comprising the nucleic acid of claim 9.
- 12. (Withdrawn) A host cell comprising the nucleic acid of claim 9.
- 13. (Withdrawn) The host cell of claim 12, which is eukaryotic or prokaryotic.
- 14. (Original) The polypeptide of claim 1, wherein the polypeptide enhances binding of low-density lipoprotein (LDL) or very low density lipoprotein (VLDL) to a cell.
- 15. (Original) The polypeptide of claim 1, wherein the polypeptide enhances degradation of low-density lipoprotein (LDL) or very low density lipoprotein (VLDL) by a cell.
- 16. (Currently Amended) A composition comprising the polypeptide of claims 1 claim 1 and a pharmaceutically acceptable carrier.
- 17. (Original) The composition of claim 16, wherein the carrier comprises dimyristoylphosphatidyl (DMPC), phosphate buffered saline or a multivesicular liposome.
- 18. (Withdrawn) A monoclonal antibody that specifically binds to the polypeptide of claim 1.

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- 19. (Withdrawn) A method for enhancing LDL binding to a cell, the method comprising contacting the cell with the polypeptide of claim 1.
- 20. (Withdrawn) A method for enhancing LDL and VLDL binding to a cell in a subject, the method comprising administering the polypeptides of claim 1, or a composition thereof, to the subject in an amount effective to increase LDL and VLDL binding to the cell of the subject.
- 21. (Withdrawn) A method for reducing serum cholesterol in a subject, the method comprising the step of administering to the subject an amount of the polypeptides of claim 1, or a composition thereof, effective to increase binding of LDL and/or VLDL to cells in the subject, thereby reducing serum cholesterol in the subject.
- 22. (Withdrawn) A method for treating a subject with coronary artery disease, the method comprising the step of administering to the subject an amount of the polypeptides of claim 1, or a composition thereof, to thereby treat the subject.
- 23. (Withdrawn) A method for treating a subject with dysbetalipoproteinemia, the method comprising the step of administering to the subject an amount of the polypeptide of claim 1, or a composition thereof, to thereby treat the subject.
- 24. (Withdrawn) A method for reducing the risk of myocardial infarction in a subject, the method comprising the step of administering to the subject an amount of the polypeptide of claim 1, or a composition thereof, to thereby treat the subject.
- 25. (Withdrawn) A method for treating atherosclerosis in a subject, the method comprising the step of administering to the subject the polypeptide of claim 1, or a composition thereof.

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- 26. (Withdrawn) A recombinant cell comprising the nucleic acid of claim 9.
- 27. (Withdrawn) A recombinant cell producing the polypeptide of claim 1.
- 28. (Withdrawn) A transgenic, non-human subject comprising the nucleic acid of claim 9.
- 29. (Withdrawn) The transgenic subject of claim 28, wherein the subject is an animal or a plant.
- 30. (Withdrawn) A transgenic non-human subject expressing the polypeptide of claim1.
- 31. (Withdrawn) The method of claim 19, wherein the administration is oral, parenteral, by intramuscular injection, by intraperitoneal injection, transdermal, extracorporeal, topical, intranasal or by inhalant.
- 32. (Withdrawn) The method of claim 19, wherein the subject is a human subject.
- 33. (Withdrawn) The method of claim 19, wherein the subject is mammal is a mouse, a rat, a rabbit, a cow, a sheep, a pig, or a primate.
- 34. (Withdrawn) The method of claim 33, wherein the primate is a human, a monkey, an ape, a chimpanzee, or an orangutan.

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